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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.	
09/996,128	11/27/2001	Alan N. Houghton	MSK.P-026-3	3698	
21121 7	590 06/25/2004		EXAM	EXAMINER	
OPPEDAHL AND LARSON LLP			HARRIS, ALANA M		
P O BOX 5068 DILLON, CO 80435-5068			ART UNIT	PAPER NUMBER	
DILLON, CO	00433 3000		1642		
			DATE MAILED: 06/25/2004		

Please find below and/or attached an Office communication concerning this application or proceeding.

	Application No.	Applicant(s)			
	09/996,128	HOUGHTON ET AL.			
Office Action Summary	Examiner	Art Unit			
	Alana M. Harris, Ph.D.	1642			
The MAILING DATE of this communication app Period for Reply	ears on the cover sheet with the c	orrespondence address			
A SHORTENED STATUTORY PERIOD FOR REPLY THE MAILING DATE OF THIS COMMUNICATION. - Extensions of time may be available under the provisions of 37 CFR 1.13 after SIX (6) MONTHS from the mailing date of this communication. - If the period for reply specified above is less than thirty (30) days, a reply - If NO period for reply is specified above, the maximum statutory period w - Failure to reply within the set or extended period for reply will, by statute, Any reply received by the Office later than three months after the mailing earned patent term adjustment. See 37 CFR 1.704(b).	36(a). In no event, however, may a reply be time within the statutory minimum of thirty (30) days fill apply and will expire SIX (6) MONTHS from cause the application to become ABANDONE!	nely filed s will be considered timely. the mailing date of this communication. D (35 U.S.C. § 133).			
Status					
1) Responsive to communication(s) filed on					
,—					
3) Since this application is in condition for allowance except for formal matters, prosecution as to the merits is closed in accordance with the practice under <i>Ex parte Quayle</i> , 1935 C.D. 11, 453 O.G. 213.					
closed in accordance with the practice under E	х рапе Quayle, 1935 С.D. 11, 45	03 O.G. 213.			
Disposition of Claims					
4) Claim(s) 1-27 is/are pending in the application. 4a) Of the above claim(s) is/are withdraw 5) Claim(s) is/are allowed. 6) Claim(s) is/are rejected. 7) Claim(s) is/are objected to. 8) Claim(s) 1-27 are subject to restriction and/or expressions.	vn from consideration.				
Application Papers					
9) The specification is objected to by the Examine 10) The drawing(s) filed on is/are: a) accomplished any objection to the	epted or b)□ objected to by the I				
Replacement drawing sheet(s) including the correct					
11) The oath or declaration is objected to by the Ex					
Priority under 35 U.S.C. § 119		`# 			
12) Acknowledgment is made of a claim for foreign a) All b) Some * c) None of: 1. Certified copies of the priority documents 2. Certified copies of the priority documents 3. Copies of the certified copies of the priority documents application from the International Bureau * See the attached detailed Office action for a list	s have been received. s have been received in Applicati ity documents have been receive u (PCT Rule 17.2(a)).	on No ed in this National Stage			
Attachment(s)	"D.,	(DTO 442)			
 Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review (PTO-948) Information Disclosure Statement(s) (PTO-1449 or PTO/SB/08) Paper No(s)/Mail Date 	4) Interview Summary Paper No(s)/Mail Da 5) Notice of Informal F 6) Other:				

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Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C.
 121:

- I. Claims 1, 2, 4, 5, 10-12, 17 and 19-24, drawn to a method for treating melanoma in a mammalian subject, comprising the step of administering to the subject an immunologically-effective amount of a xenogeneic differentiation antigen, wherein said antigen is human tyrosinase, classified in class 514, subclass 8. Claims 1, 4, 10, 11, 17, 20, 23 and 24 will be examined with this Group to the extent that the xenogeneic differentiation antigen is a human tyrosinase.
- II. Claims 1, 3, 4, 6, 10, 11, 13, 19, 20, 23 and 24, drawn to a method for treating melanoma in a mammalian subject, comprising the step of administering to the subject an immunologically-effective amount of a xenogeneic differentiation antigen, wherein said antigen is human gp75, classified in class 436, subclass 64. Claims 1, 4, 10, 11, 19, 20, 23 and 24 will be examined with this Group to the extent that the xenogeneic differentiation antigen is a human gp75.
- III. Claims 1, 7-11, 14, 15, 18-20, 23 and 24, drawn to a method for treating melanoma in a mammalian subject, comprising the step of administering to the subject an immunologically-effective amount of a xenogeneic differentiation antigen, wherein said antigen is a murine tyrosinase, classified in class 424, subclass 184.1. Claims 1, 4, 7-10, 11, 14, 19, 20, 23 and 24 will be examined with this

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Group to the extent that the xenogeneic differentiation antigen is a murine tyrosinase.

- IV. Claims 1, 7-10, 14, 16, 19, 20, 23 and 24, drawn to a method for treating melanoma in a mammalian subject, comprising the step of administering to the subject an immunologically-effective amount of a xenogeneic differentiation antigen, wherein said antigen is murine gp75, classified in class 424, subclass 184.1. Claims 1, 7-10, 14, 19, 20, 23 and 24 will be examined with this Group to the extent that the xenogeneic differentiation antigen is a murine gp75.
- V. Claims 26, drawn to a vector comprising SEQ ID NO: 1, classified in class 435, subclass 320.1.
- VI. Claim 27, drawn to a vector comprising SEQ ID NO: 2, classified in class 435, subclass 320.1.
- 2. The inventions are distinct, each from the other because of the following reasons:

Inventions I and V are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case the vector of Group V, which encodes the human tyrosinase

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implemented in the *in vivo* method of Group I could also be used in an *in vitro* reporter assay.

Inventions II and VI are related as product and process of use. The inventions can be shown to be distinct if either or both of the following can be shown: (1) the process for using the product as claimed can be practiced with another materially different product or (2) the product as claimed can be used in a materially different process of using that product (MPEP § 806.05(h)). In the instant case the vector of Group VI, which encodes the murine tyrosinase implemented in the *in vivo* method of Group II could also be used in an *in vitro* reporter assay.

Inventions I-IV are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the different inventions of all four groups read on a method of treating melanoma comprising administering a xenogeneic differentiation antigen. However, each differentiation antigen is patentably distinct and structurally different and would elicit a different immune response. Groups I-IV use the differentiation antigens, human tyrosinase, human gp75, murine tyrosinase and murine gp75, respectively.

Inventions V and VI are unrelated. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the different inventions of Groups V

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and VI both are both vectors, which comprise polynucleotide sequences, however the sequences are distinct and encode structurally different proteins.

- 3. Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their different classification, restriction for examination purposes as indicated is proper.
- 4. A telephone call was made to Marina Larson on June 24, 2004 to request an oral election to the above restriction requirement, but did not result in an election being made.
- 5. Applicant is advised that the reply to this requirement to be complete must include an election of the invention to be examined even though the requirement be traversed (37 CFR 1.143).
- 6. Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

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7. Any inquiry concerning this communication or earlier communications from the examiner should be directed to Alana M. Harris, Ph.D. whose telephone number is (571)272-0831. The examiner works a flexible schedule, but can

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Christine Y. Chan can be reached on (703)272-0841. The fax phone number for the organization where this application or proceeding is assigned is 703-872-9306.

normally be reached between the hours of 6:30 am to 5:00 pm.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

ALANA M. HARRIS, PH.D. PRIMARY EXAMINER

Alana M. Harris, Ph.D.

17 June 2004